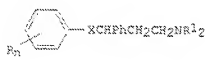
AN 1977:405817 CAPLUS
DN 87:5817
TI Piperidinopropyl phenyl ethers
IN Ogawa, Shuntaro; Morita, Minoru; Yoshida, Akiyoshi
PA Rohto Pharmaceutical Co., Ltd., Japan
SO Japan., 10 pp.
CODEN: JAXXAD
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 52000941	B4	19770111	JP 1969-99282	19691209

GI



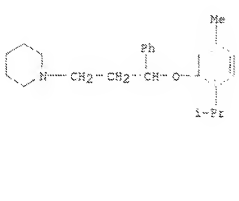
I

AB About 16 piperidinopropyl phenyl ethers I (Rn = H, 2-, 4-Me, 4-EtO, 2-Me2CH-4-Me; NR12 = NMe2, NMe2, piperidino, 2- or 4-methylpiperidino, pyrrolidino; X = O, S) were prepd. from RnC6H5-nOH with ClCHCH2CH2NR12. Thus, 2.7¼ g 1-chloro-1-phenyl-3-piperidinopropane was refluxed 0.94 g PhOH in Me2CHOH contg. Na to give 2.¢ g I (Rn = H, NR12 = piperidino, X = O). ED50 values for their anticonvulsant activity are tabulated.

IT 62896-92-8P
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 62896-92-8 CAPLUS

CN Piperidine, 1-[3-[5-methyl-2-(1-methylethyl)phenoxy]-3-phenylpropyl]- (9CI) (CA INDEX NAME)



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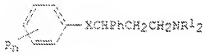

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IT 62663-41-6P
RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(prepn. and anticonvulsant activity of)

RN 62663-41-6 CAPLUS
CN Piperidine, 1-[3-[(4-methylphenyl)thio]-3-phenylpropyl]- (9CI) (CA INDEX NAME)